Sun-Shine Land International Ltd.
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EU Declaration of Conformity

According to the Medical Device Regulation: 2017/745/EU

Manufacturer

Sun-Shine Land International Ltd.

8F, No. 69-10, Sec. 2, Chung Cheng E. Rd., Tamsui Dist., New Taipei City, Taiwan,

Authorized representative in Europe

Y. Sung Handelsvertretung

Duesselthaler Str. 24, 40211 Duesseldorf, Germany

We declare that the products described hereafter:

Model	Description
BA-7242	Bedpan
BA-7246	Bedpan with Lid
BA-7247	Male Urinal w/ Lid
BA-7248	Female Urinal
HA-4770	Transfer Turntable

that are covered by the present declaration is in conformity with the Medical Device **Regulation 2017/745/EU**. The device is in conformity with conformity assessment procedure for **Class I devices** that should be carried out, as a general rule, under the sole responsibility of manufacturers in view of the low level of vulnerability associated with such devices. Thus, manufacturers of class I devices, other than custom-made or investigational devices, shall declare the conformity of their products by issuing an EU declaration of conformity referred to in Article 19 "EC declaration of conformity" after drawing up the technical documentation set out in Annexes II and III of the **Regulation**.

Authority: David Yu/ Manager
December 23, 2021